CLAIMS

1. The peptides, being either epitopes or potential epitopes for the stated HLA (human leucocyte antigen) class I molecules, conservative variants thereof, and longer peptides containing these sequences which are sub-units of the indicated antigens:

label					Se	drevce	1				Position
HLA-A2	1	2	3	4	5	6	7	8	9	10	
tr26	H	L	G	N	v	K	Y	L	v		3
tr29	L	L	M	D	C	s	G	s	I		51
tr39	G	I	λ	G	G	L	λ	L	L		500
_# ls10	ı	L	Y	I	s	F	Y	F	I		4
ls1 1	Y	I	s	F	Y	F	I	L	v		6 .
1819	G	I	Y	K	E	L	E	D	L		1801
1 823	H	I	F	D	G	D	N	E	I		1883
Ep36	¥	L	K	T	I	Q	n	8	L		334
cp37	Y	L	Q	ĸ	I	Q	N	s	L		334
cp38	Y	L	Q	K	I	K	n	s	L		334
cp39	Y	L	N	ĸ	I	Q	N	s	L		334
HLA-B8							**				
cp43	L	R	K	P	Ř,	H	, x	K	L		134
cp44	L	ĸ	K	I	K	N	S	I	s		335
cp45	Q	v	R	I	ĸ	P	G	s	A		358
cp46	A	N	K	P	K	D	G	L	D		366
tr42	λ	s	ĸ	N	K	K	ĸ	λ	L		107
tr43	ĸ	N	ĸ	R	ĸ	λ	L	I	I		109

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	label					Seq	ience		•		Po	sition
	•	1	2	3	4	5	6	7 ·	8	9	10	
	HLA-B17											
	cp48	ŗ	s	v	S	s	F	L	F	v		8
	cp55	G	S	A	N	K	P	K	D .	E	L	364
	cp56	С	s ·	S	v	F	N	v	v			388
	ls36	N	s	E	K	D	E	I	I			28
	ls37	G	s	S	N	S	R	N	R	I		42
	ls39	v	S	Q	T	N	F	K	s	L		92
	ls40	K	s	L	L	R	N	L	G	v		98
ļ.	ls42	Q	s	D	s	E	Q	E	R	L		179
4000	ls45	R	T	K	A	S	ĸ	E	T	L		1187
) 5	ls48	H	T	L	E	T	v	N	I			1742
	1s49	I	S	D	v	N	D	F	Q	I		1749
K	1s50	I	s	K	Y	E	D	E	I			1757
ř	ls51	ı	S	A	E	Y	D	D	s	L		1764
i.	1853	K	S	L	Y	D	E	H	I			1854
:	1s54	L	S	E	D	I	T	ĸ	Y	F		1898
	ls55	T	K	Y	F	M	K	L				1902
ļ	tr57	K	T	A	S	С	G	V	W	D	EW	240
į	tr58	G	T	R	S	R	K	R	B	I	L	260
	tr59	S	S	V	Q	K	P	E	E	N	I	311
	tr60	D	S	E	K	E	v	P	s	D	v	367
	tr61	Y	S	P	L	P	P	K	v	L		415
	tr62	E	S	D	N	ĸ	Y	K	I	A		490
	tr63	A	T	P	Y	A	G	E	P	A		523
	tr64	E	T	L	G	E	E	D	K	D	L	535

these peptides being selected from three *Plasmodium* falciparum antigens, circumsporozoite protein (cp), thrombospondin-related anonymous protein (tr) and liver-stage antigen-1 (ls),

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- 2. A peptide comprising at least two of the sequences listed in claim 1.
- A peptide as claimed in claim 1 or claim 2 having an N-terminus or C-terminus carrying a lipid tail.
- 4. A peptide as claimed in any one of claims 1 to 3, comprising 8-100 amino acid residues.
- 5. A vaccine comprising at least one peptide according to any one of claims 1 to 4, for immunisation against malaria.
- 6. Use of *Plasmodium falciparum* gene or protein TRAP (thrombospondin-related anonymous protein) as a cytotoxic T lymphocyte-inducing gene or protein for immunization against malaria.
- 7. Oligonucleotides which code for the peptides claimed in any one of claims 1 to 4.
 - 8. A vaccine comprising at least one oligonucleotide according to claim 7 for expression *in vivo* for immunization against malaria.
- 20 9. A method of inducing primary cytotoxic T lymphocyte responses to a chosen antigen or microorganism, which method comprises incubating lymphocytes ex vivo with the chosen antigen or microorganism in the presence of KLH (keyhole limpet
- haemocyanin) or any other substance which preferentially stimulates a CD45RA+ subset of T lymphocyte.
- 10. A method as claimed in claim 9, wherein IL-7 (interleukin-7) and/or IL-2 (interleukin-2) is also present during incubation.

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11.	IIca	οf	2017	One	Ωf	the	peptides	
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label	Sequence										Position
HLA-B7	1	2	3	4	5	6	7	8	9	10	
cp6	, M	P	N	D	P	N	R	n	v		300
cp6.1	M	P	×	Y	P	N	R	N	v		300
cp6.2	ĸ	P	N	N	P	N	R	N	v		300
ls6	ĸ	P	I	٧	Q	Y	D	N	F		1786
shl	I	P	s	L	A	L	M	L	I		7
sh6	M	P	L	E	T	Q	L	A	I		77
cp21	N	P	D	P	N	A	N	P	N	v	120
tr6	N	P	E	N	P	P	N	P	D	I	348
tr13	I	P	D	s	I	Q	D	s	L		164
tr15	E	P	A	P	F	D	E	T	L		529
tr21	G	P	F	M	K	A	v	C	V		228

and conservative variants thereof and longer peptides containing the sequences which are sub-units of the stated antigen, and of oligonucleotides which code for the said peptides, as a cytotoxic T lymphocyte-inducer for immunization against malaria of individuals possessing a HLA-B7 allele.